

Fr m: STIC-ILL
Sent: Wednesday, March 06, 2002 9:19 AM
T : STIC-Biotech/ChemLib
Subject: FW: sequence search request

-----Original Message-----

From: Chernyshev, Olga
Sent: Wednesday, March 06, 2002 9:17 AM
To: STIC-ILL
Subject: sequence search request

US case 09/853,753
Please search SEQ ID NO:1 and 2.
Thank you very much!

Olga N. Chernyshev
AU1646
CM1 8D06
305-1003
mail box 10C01

Edward Hart
Technical Info. Specialist
STIC/Biotech
CMI 6B02 Tel: 305-9203

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Searcher: _____
Phone: _____
Location: _____
Date Picked Up: 3/7/02
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Searcher Prep/Review: _____
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Online time: _____

TYPE OF SEARCH:
NA Sequences: _____
AA Sequences: _____
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST(where applic.)
STN: _____
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Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
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WWW/Internet: _____
Other (specify): _____

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	2297	100.0	2397	9	AF254868	AF254868 Homo sapi
2	2241	97.6	2713	9	HS4278865	AI2728865 Homo sapi
3	2163	94.2	133568	9	HS156915	Z93015 Human DNA s
4	186	8.1	1818	9	SS83462	S83462 ALS-85 kda
5	183.2	8.0	37635	9	AC0004152	AC0004152 Homo sapi
6	173.4	7.5	2125	9	HOMOIGF1	M86882 Human IGF f
7	173.4	7.5	4574	9	AF192554	AF192554 Homo sapi
8	173.4	7.5	21419	9	HS44786	AI031724 Human DNA
9	161.2	7.0	251997	2	AC012180	AC012180 Homo sapi
10	147.8	6.4	128318	2	AC0925332	AC0925332 Papio cyn
11	145.6	6.3	157419	2	AC025389	AC025389 Homo sapi
12	144.6	6.3	1907	9	AB045987	AB045987 Macaca fa
13	143	6.2	1441	9	AF283463	AF283463 Homo sapi
14	143	6.2	1782	9	BC011787	BC011787 Homo sapi
15	143	6.2	2236	6	AX047642	AX047642 Sequence
16	143	6.2	2236	6	AX055398	AX055398 Sequence
17	143	6.2	168339	9	AC007663	AC007663 Homo sapi
18	143	6.2	174840	9	AC006549	AC006549 Homo sapi
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20	141.8	6.2	2190	10	S46785	S46785 insulin-I like
21	140.2	6.1	5270	10	AF006203	AF006203 Rattus nor
22	137.8	5.0	154277	2	AP003943	AP003943 Oryza sat
23	131.4	5.7	2621	9	AB046639	AB046639 Macaca fa
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25	131.2	5.7	2636	6	AX156560	AX156560 Sequence
26	130.4	5.7	215727	2	AC073701	AC073701 Mus muscu
27	130.2	5.7	4933	9	AB014544	AB014544 Homo sapi
28	129.8	5.7	2257	6	AX083306	AX083306 Sequence
29	129.8	5.7	129.8	6	BC011057	BC011057 Homo sapi
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31	128.6	5.6	195910	9	AC005013	AC005013 Homo sapi
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33	127.6	5.6	2862	9	HDMCRAN	J05188 Human carbo
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35	126	5.5	175759	10	AF220294	AF220294 Mus muscu
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37	125.2	5.5	159802	2	AC019262	AC019262 Homo sapi
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39	124	5.4	136198	2	AP003883	AP003883 Oryza sat
40	122.4	5.3	251997	2	AC012180	AC012180 Homo sapi
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RESULT	1
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DEFINITION	Homo sapiens nycaloplin mRNA, complete cds.
ACCESSION	AF254868
VERSION	AF254868.1 GI:11993320
KEYWORDS	human.
SOURCE	Homo sapiens
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo. 1 (bases 1 to 2297)
REFERENCE	Bech-Hansen,N.T., Naylor,M.J., Maybaum,T.A., Sparkes,R.L., Koop,B. Dirck,D.G., Bergen,A.A., Prinsen,C.F., Polomeno,R.C., Gal,A., Birch,A.V., Musarella,M.A., Jacobson,S.G., Young,R.S. and Melcher,R.C. Mutations in NYX, encoding the leucine-rich proteoglycan nycaloplin, cause X-linked complete congenital stationary night blindness Nat. Genet. 26 (3), 319-322 (2000)
TITLE	
JOURNAL	

MEDLINE	20517340
PUBMED	11062471
REFERENCE	2 (bases 1 to 2297)
AUTHORS	Bech-Hansen, N.T., Maybaum, T.A. and Naylor, M.J.
TITLE	Direct Submission
JOURNAL	Submitted (10-APR-2000) Medical Genetics, University of Calgary
FEATURES	3330 Hospital Dr. N.W., Calgary, AB T2N 4N1, Canada
	Location/Qualifiers

CDS

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Query Match	100.0%	Score 2297	DB 9	Length 2297
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[illegible]

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RESULT 2
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LOCUS
DEFINITION HSA278865 2713 bp mRNA
ACCESSION AJ278865
VERSION AJ278865.1 GI:11877211
KEYWORDS CLRP gene; leucine-rich repeat protein.
SOURCE human.
ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE
AUTHORS Pusch, C.M., Zeltz, C., Brandau, O., Pesch, R., Achatz, H., Fell, S.,
 Schafke, C., Maurer, J., Jacobi, F.K., Pluckers, A., Andreasson, S.,
 Hardcastle, A., Wessinger, B., Berger, W. and Weidl, A.
TITLE The complete form of X-linked congenital stationary night blindness
 is caused by mutations in a gene encoding a leucine-rich repeat
 protein
JOURNAL Nat. Genet. 26 (3), 324-327 (2000)
MEDLINE 20517341
REFERENCE 2 (bases 1 to 2713)
AUTHORS Pusch, C.M.
TITLE Direct Submission
JOURNAL Submitted (23-AUG-2000) Medical Genetics, Ludwig's Maximilians
 University, Goethestr. 29, Munich D-80336, GERMANY
FEATURES
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